In the claims:

1. (Currently Amended) A pharmaceutical composition for intramammary administration to a non-human mammal, comprising an antibacterial agent, prednisolone and a pharmaceutically acceptable carrier, ~~characterised in that~~ wherein the composition comprises at least 20 mg of prednisolone / unit dose.

2. (Currently Amended) The composition according to claim 1, ~~characterised in that it comprises~~ comprising the prednisolone in an amount of 20 to 40 mg / unit dose.

3. (Currently Amended) The composition according to claim 2, ~~characterised in that it comprises~~ comprising the prednisolone in an amount of 20 to 30 mg / unit dose.

4. (Currently Amended) The composition according to ~~any of claims 1 to 3, characterised in that~~ claim 1, wherein the antibacterial agent is a cephalosporin.

5. (Currently Amended) The composition according to claim 4, ~~characterised in that~~ wherein the cephalosporin is cephapirin.

6. (Currently Amended) The composition according to claim 4, ~~characterised in that~~ wherein the cephalosporin is cefquinome.

7. (Currently Amended) The composition according to ~~any of claims 1 to 6, characterised in that it comprises~~ claim 1, comprising the antibacterial agent in an amount of 10 to 500 mg/ unit dose.

8. (Currently Amended) A process for preparing a pharmaceutical composition ~~as claimed in any of claims 1 to 7~~ according to claim 1, comprising the steps of mixing an oil and one or more ~~optionally~~ pharmaceutically acceptable additives to form a carrier, and suspending the antibacterial agent and the prednisolone in the carrier.

9. (Canceled).